

K222780 LILY Extension Tube and Needleless ConnectorNov 9, 2023
421 days to decisionK222780 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k222780/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Sep 14, 2022
Decision date	Nov 9, 2023
Days to decision	421 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Lily Medical Corporation
Location	Miaoli County, CN
Contact	Steven Shen
510(k) history	1 submissions · 1 cleared · 2023-2023

REGULATORY CONSULTANT

Consulting firm	Benq Medical Technology Corporation
Contact	Steven Shen

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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